STN Search History

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PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

NEWS 1 Web Page for STN Seminar Schedule - N. America

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PASSWORD:

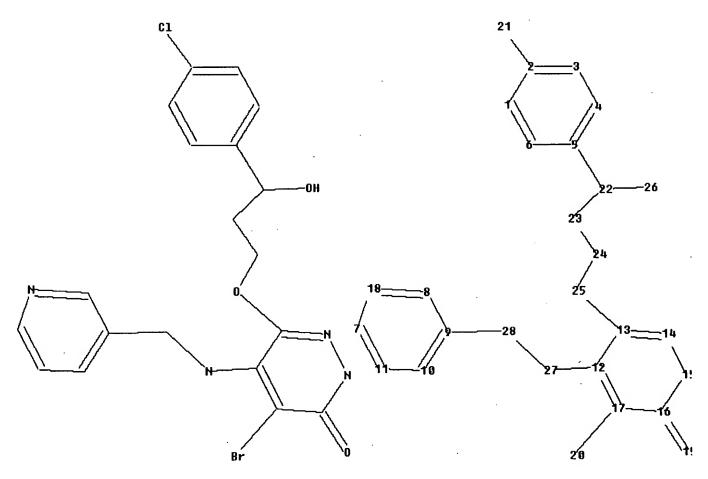
\* \* \* \* \* RECONNECTED TO STN INTERNATIONAL \* \* \* \* \* \* \* \* SESSION RESUMED IN FILE 'REGISTRY' AT 17:01:05 ON 17 SEP 2007 FILE 'REGISTRY' ENTERED AT 17:01:05 ON 17 SEP 2007 COPYRIGHT (C) 2007 American Chemical Society (ACS)

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 173.45 384.30

FULL ESTIMATED COST

Uploading C:\Program Files\Stnexp\Queries\10584222-1.str



chain nodes :
19 20 21 22 23 24 25 26 27 28
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18
chain bonds :
2-21 5-22 9-28 12-27 13-25 16-19 17-20 22-23 22-26 23-24 24-25 27-28
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-11 7-18 8-9 8-18 9-10 10-11 12-13 12-17 13-14 14-15 15-16 16-17
exact/norm bonds :
12-13 12-17 13-14 14-15 15-16 16-17 16-19
exact bonds :
2-21 5-22 9-28 12-27 13-25 17-20 22-23 22-26 23-24 24-25 27-28
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-11 7-18 8-9 8-18 9-10 10-11

## Match level :

L8

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS

## STRUCTURE UPLOADED

=> s 18
SAMPLE SEARCH INITIATED 17:01:41 FILE 'REGISTRY'

100.0% PROCESSED

1 ITERATIONS

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:

ONLINE \*\*COMPLETE\*\*

BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS:

1 TO 80

PROJECTED ANSWERS:

1 TO.

L9

80

1 SEA SSS SAM L8

=> d 19

ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN L9

171661-81-7 REGISTRY RN

ED Entered STN: 22 Dec 1995

3(2H)-Pyridazinone, 4-bromo-6-[3-(4-chlorophenyl)-3-hydroxypropoxy]-5-[(3-CN pyridinylmethyl)amino]-, (S)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

C19 H18 Br Cl N4 O3 MF

COM CI

SR CA

LC

STN Files: CA, CAPLUS, CASREACT, USPATFULL

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> fil caplus medline

SINCE FILE

TOTAL

COST IN U.S. DOLLARS

ENTRY

SESSION

FULL ESTIMATED COST

177.65 388.50

FILE 'CAPLUS' ENTERED AT 17:04:03 ON 17 SEP 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'MEDLINE' ENTERED AT 17:04:03 ON 17 SEP 2007

L10 1 L9

=> d l19 ibib abs hitstr

L19 NOT FOUND

The L-number entered has not been defined in this session, or it has been deleted. To see the L-numbers currently defined in this session, enter DISPLAY HISTORY at an arrow prompt (=>).

## => d l10 ibib abs hitstr

L10 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 1995:992456 CAPLUS Full-text

DOCUMENT NUMBER: 124:55968

TITLE: Preparation of pyridazinone derivatives having potent

antithrombocytic activity

INVENTOR(S): Tanikawa, Keizo; Matsumoto, Takashi; Matsumoto, Hiroo;

Tsuruzoe, Nobutomo; Nakabeppu, Hitoshi

PATENT ASSIGNEE(S): Nissan Chemical Industries, Ltd., Japan

SOURCE: PCT Int. Appl., 32 pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Facence English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

						APPLICATION NO. DATE	DATE	
						WO 1995-JP69 19950124		
	W: AU,	CA,	CN,	CZ,	FI, HU, KR,	MX, NO, NZ, RO, RU, SI, SK, UA, US		
	RW: AT,	BE,	CH,	DE,	DK, ES, FR,	GB, GR, IE, IT, LU, MC, NL, PT, SE		
CA	2181901			A1	19950727	CA 1995-2181901 19950124 AU 1995-14663 19950124		
CA	2181901			С	20050913			
ΑU	9514663			Α	19950808	AU 1995-14663 19950124		
EP	742211			A1	19961113	EP 1995-906505 19950124		
					20000510			
	R: AT,	BE,	CH,	DE,	DK, ES, FR,	GB, IT, LI, NL, PT, SE		
CN	1138852			Α	19961225	CN 1995-191304 19950124		
CN	1049892			В	20000301	CN 1995-191304 19950124		
HII	74742			Δ2	19970228	ни 1996-2021 19950124		
HU	223963			B1	20050329			
AT	192741			T	20000515	AT 1995-906505 19950124		
ES	2147841			Т3	20001001	ES 1995-906505 19950124		
PT	742211			T	20001031	AT 1995-906505 19950124 ES 1995-906505 19950124 PT 1995-906505 19950124		
JР	07252237	•		Α	19951003	JP 1995-9398 19950125		
JP	3666042			B2	20050629			
TW	420665			В	20010201	TW 1995-84100797 19950127		
US	5750523			Α	19980512	TW 1995-84100797 19950127 US 1996-676227 19960723		
					19960724			
	112214			В1				
NO	9603095			Α	19960924	NO 1996-3095 19960724		
NO	307965			В1	20000626			
	5856327					US 1997-936600 19970924	;	
	Y APPLN.					JP 1994-6541 A 19940125		
			-			WO 1995-JP69 W 19950124		
						US 1996-676227 A3 19960723		

OTHER SOURCE(S): CASREACT 124:55968; MARPAT 124:55968

GI

Pyridazinone derivs. represented by general formula [I; R = H, Cl-4 alkyl; X = AB H, Cl, Br; Ar = pyridyl, Ph substituted by OR1 (wherein R1 = H or Cl-4 alkyl) and a group selected from H, halo, or C1-4 alkyl or a group selected from OH or C1-4 alkoxy; Y = C1-8 alkylene, one of its C atom being substituted by one OR1 group; Z1, Z2 = H, halo, C1-4 alkyl, OR1 (R1 being as defined above)], which have a broad spectrum of blood platelet aggregation inhibition with high selectivity and reduced side effects (e.g. headache, heaviness of head, hypotension, and palpitation) and are safely used as the active ingredient of a preventive or remedy for various thrombotic diseases, are prepared Thus, a mixture of 1.50 g 4,5-dibromo-6-[3-(4-chlorophenyl)-3-hydroxypropyloxy]-3(2H)pyridazinone, 1.48 g 3-picolylamine, 45 mL MeOH, and 5 mL H2O was refluxed with stirring overnight to give 1.05 g of the title compound (II; R2 = OH). This compound in vitro inhibited the ADP- and collagen-induced blood platelet aggregation of rabbit platelet rich plasma with IC50 of 0.23 and 0.099  $\mu\text{M},$ resp. It in vitro showed weaker vasodilating activity (EC50 of 1.3  $\mu M$ ) than the known compound II.HCl (R2 = H) (EC50 of 0.4  $\mu M$ ) in an assay of inhibiting the phenylephrine-induced contraction of rabbit thoracic aorta rings. A tablet and a capsule formulation containing II (R2 = OH) were described. IT 171661-81-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyridazinone derivs. having potent antithrombocytic activity)

RN 171661-81-7 CAPLUS

CN 3(2H)-Pyridazinone, 4-bromo-6-[3-(4-chlorophenyl)-3-hydroxypropoxy]-5-[(3-pyridinylmethyl)amino]-, (S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

=> logoff hold SINCE FILE TOTAL COST IN U.S. DOLLARS ENTRY SESSION FULL ESTIMATED COST 9.28 397.78 SINCE FILE TOTAL DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) ENTRY SESSION -0.78 -0.78

SESSION WILL BE HELD FOR 120 MINUTES STN INTERNATIONAL SESSION SUSPENDED AT 17:09:00 ON 17 SEP 2007

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SEP 17

patents

NEWS 22

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NEWS EXPRESS 05 SEPTEMBER 2007: CURRENT WINDOWS VERSION IS V8.2, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 05 SEPTEMBER 2007.

CAplus coverage extended to include traditional medicine